

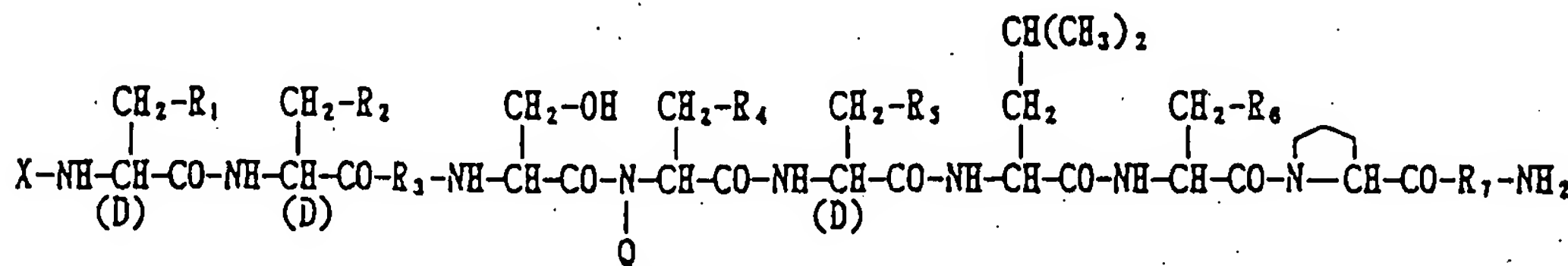
**IN THE ABSTRACT:**

Please substitute the attached page with a new Abstract in place of the originally filed Abstract.

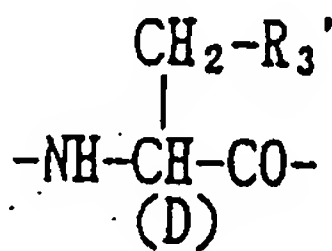
**IN THE CLAIMS:**

**Please amend the claims as follows:**

1. (currently amended) A sustained-release preparation which comprises a physiologically active peptide of the general formula



wherein X represents an acyl group; R<sub>1</sub>, R<sub>2</sub> and R<sub>4</sub> each represents an aromatic cyclic group; R<sub>3</sub> represents a D-amino acid residue or a group of the formula



wherein  $R_3$  is a heterocyclic group;  $R_5$  represents a group of the formula  $-(CH_2)_n-R_5$ , wherein  $n$  is 2 or 3 and  $R_5$  is an amino group which ~~may~~is optionally ~~be~~ substituted, an aromatic cyclic group or an O-glycosyl group;  $R_6$  represents a group of the formula  $-(CH_2)_n-R_6$ , wherein  $n$  is 2 or 3 and  $R_6$  is an amino group which ~~may~~is optionally ~~be~~ substituted;  $R_7$  represents a D-amino acid residue or an azaglycyl residue; and  $Q$  represents hydrogen or a lower alkyl group, or a salt thereof; and a biodegradable polymer having a terminal carboxyl group.

2. (currently amended) The sustained-release preparation according to claim 1, wherein X is a C<sub>2-7</sub> alkanoyl group which ~~may~~is optionally ~~be~~-substituted by a 5- or 6-membered heterocyclic carboxamido group.

## ABSTRACT

A sustained-release preparation which comprises a physiologically active peptide of general formula I and a biodegradable polymer having a terminal carboxyl group. The sustained-release preparation shows a constant release of the peptide over a long period of time and is substantially free from an initial burst.